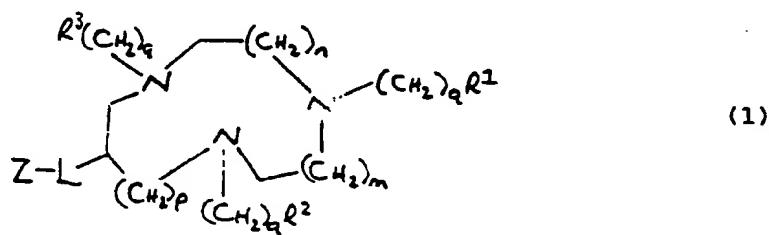




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(54) Title: TRI-AZA MACROCYCLES AND PROCESSES FOR THEIR PREPARATION



(57) Abstract

Tri-aza macrocycles of formula (1), wherein m and n, which may be the same or different, is each zero or an integer 1, 2, or 3; p is zero or an integer 1 or 2; q is zero or an integer from 1 to 3 inclusive; R¹, R² and R³, which may be the same or different, is each a hydrogen atom or a group -CO₂H or -P(O)(XH)R⁴ where X is an oxygen or sulfur atom and R⁴ is a hydrogen atom or an alkyl or alkoxy group), with the proviso that at least one of R¹, R² and R³ is a -P(O)(XH)R⁴ group; L is a covalent bond or a linker group; Z is a hydrogen atom or a reactive functional group; and metal complexes and/or salts thereof; are described together with processes for their preparation and compositions containing them. The compounds are useful as imaging agents, and in the treatment of abnormal cell disorders, such as in the treatment of tumours, and may be coupled to other molecules, such as proteins, for use in diagnosis and therapy.

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TRI-AZA MACROCYCLES AND PROCESSES FOR THEIR PREPARATIONField of the Invention

5 This invention relates to functionalised tri-aza macrocycles, to metal complexes thereof, to conjugate compounds containing the functionalised tri-aza macrocycles and metal complexes thereof, and to their use in diagnosis and therapy.

Background to the Invention

10 The attachment of metal ions to proteins, peptides and other, smaller molecules is a fast expanding technology, which has numerous proven and potential applications in research, in industry and, particularly, in medicine.

15 In recent years, much of the impetus behind the development of this technology has been the ability to link metal ions to antibodies, especially monoclonal antibodies. Such metal labelled antibodies have found a widespread use, especially in medicine, where they have been employed, for example, to target the metal ions to a specific tissue type, both in vitro and in vivo. Thus, metal labelled antibodies have applications in locating specific tissue types (e.g. employing computer-aided tomographic techniques where the metal ion is in some way detectable) and in the treatment of cell disorders (e.g. treating mammalian tumours where the metal ion is a cytotoxic radionuclide).

20 25 Conventionally, attachment of the metal ion to a protein such as an antibody has been achieved by complexation by an acyclic chelate such as a substituted diethylenetriaminepentaacetic acid [Gansow O. A. et al, Inorg. Chem., (1986), 25, 2772, and see for example, U.S. Patent No. 4454106] or ethylenediaminetetraacetic acid [Meares, C. F. et al, Acc. Chem. Res., (1984), 17, 202] covalently linked to the antibody. Such acyclic complexes however tend to be unstable in vivo either as a result of acid-catalysed decomplexation or

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5 competitive chelate binding by Ca^{2+} or Zn^{2+} in serum, or as a result of competition from transferrin [Moerlein, S. M. et al, Int. J. Nuc. Med. Biol., (1981) 8, 277]. The lack of stability can result in uncomplexed metal atoms in the body which have a cytotoxic effect on healthy tissue (e.g. bone marrow) or which markedly reduce the signal-to-noise ratio of an imaging technique.

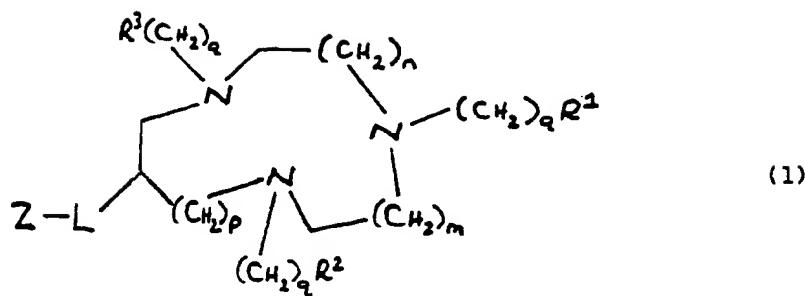
10 A possible alternative to the use of acyclic chelates in the labelling of antibodies is the use of macrocyclic ligands, which has previously been suggested in broad terms [Gansow O. A. et al, Am. Chem. Soc. Symp. Ser., (1984), 241, 215; UK Patent Specification No. 2122641; and Moi M. K. et al, Anal. Biochem., (1985), 148, 249-253]. More recently, tri-aza and tetra-aza macrocycles have been described which are capable of binding metals, and which can be conjugated to antibodies (International Patent Specifications Nos. 15 WO 89/01475 and WO89/01476).

20 Other tri-aza macrocycles have also been described, which are capable of binding metals [International Patent Specification No. WO86/02352; European Patent Specification No. 197437; Bryden, C.C. et al, Rare Earths Mod. Sci. Technol. (1982), 3, 53-57; Kabachnik, I.M. et al, Izv. Akad. Nauk. SSSR, Ser. Khim., 835 (1984)]. Some compounds of these types, when complexed with metals, have been proposed for use as contrast agents for use in diagnostic imaging.

25 We have now found a new class of functionalised tri-aza macrocycles, members of which are able to form kinetically inert complexes with metal ions which are of use in diagnosis and therapy.

Summary of the Invention

Thus according to one aspect of the present invention we provide a compound of general formula (1):



wherein

m and n, which may be the same or different, is each zero or an integer 1, 2, or 3;

p is zero or an integer 1 or 2;

5 q is zero or an integer from 1 to 6 inclusive;

R^1 , R^2 and R^3 , which may be the same or different, is each a hydrogen atom or a group $-CO_2H$ or $-P(O)(XH)R^4$ (where X is an oxygen or sulphur atom and R^4 is a hydrogen atom or an alkyl or 10 alkoxy group), with the proviso that at least one of R^1 , R^2 and R^3 is a $-P(O)(XH)R^4$ group;

L is a covalent bond or a linker group;

Z is a hydrogen atom or a reactive functional group;

and metal complexes and/or salts thereof.

In the compounds of formula (1), alkyl groups represented by R^4 15 may be for example C_{1-6} alkyl groups such as methyl or ethyl groups.

Alkoxy groups represented by R^4 may be C_{1-6} alkoxy groups such as methoxy or ethoxy groups.

In general, compounds of formula (1) in which R^1 , R^2 and R^3 are the same and is each a group $-P(O)(XH)R^4$ are preferred.

20 Compounds of this type in which q is an integer from 1 to 6 inclusive, particularly an integer 1, are especially preferred.

Particularly useful compounds of formula (1) are those wherein R^1 , R^2 and R^3 is each a group $-P(O)(OH)H$, $-P(O)(OH)OCH_3$,

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$-P(O)(OH)OCH_2CH_3$ or especially $-P(O)(OH)R^4$ where R^4 is an alkyl group, particularly a methyl group. In compounds of this type, q is preferably an integer from 1 to 6 inclusive, particularly an integer 1.

5 In the compounds of formula (1), it will be appreciated that the nature of the group L when it is a linker group may be varied widely without substantially affecting the usefulness of compounds of formula (1) and the metal complexes thereof. Thus L may be any suitable organic radical and may be for example an optionally substituted aliphatic hydrocarbyl chain, optionally interrupted by

10 one or more heteroatoms selected from -O- or -S- or by one or more $-N(R^5)-$ (where R^5 is a hydrogen atom or a C_{1-6} alkyl group), $-CON(R^5)-$, $-N(R^5)CO-$, cycloaliphatic, aromatic, or heteroaromatic groups.

15 In the above definition, and in the same context whenever it appears below, the term "interrupted by" as applied to cycloaliphatic or aromatic groups is to be understood to also mean that these particular groups may additionally be present linked to the terminal carbon atom of the hydrocarbyl chain represented by L, at the 20 opposite end of the chain to the carbon atom attached to the macrocycle.

Thus, for example, L may be an optionally substituted straight or branched C_{1-20} alkylene, C_{2-20} alkenylene, or C_{2-20} alkynylene chain, optionally interrupted by one or more -O- or -S- atoms or 25 C_{5-8} cycloalkylene (e.g. cyclopentylene or cyclohexylene), C_{6-12} aromatic (e.g. phenylene or substituted phenylene), C_{5-10} heteroaromatic (e.g. furanyl, pyridyl), $-N(R^5)-$, $-CON(R^5)-$ or $-N(R^5)CO-$ groups.

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Examples of substituents which may be present on the chain L include halogen atoms, e.g. fluorine, chlorine, bromine, or iodine atoms or groups selected from C_{1-6} alkoxy (e.g. methoxy or ethoxy), hydroxy, nitro, $-N(R^6)(R^7)$, [where R^6 is a hydrogen atom or a

5 C_{1-6} alkyl group and R^7 is a C_{1-6} alkyl group; e.g. $-NHCH_3$ or $-N(CH_3)_2$], or substituted amido, e.g. a group of formula $-(CH_2)_nCON(R^8)(R^9)$ [where n is zero or an integer 1 to 4 inclusive, R^8 is a hydrogen atom or a C_{1-6} alkyl group, e.g. methyl and R^9 is an optionally substituted C_{1-6} alkyl group].

10 Substituted alkyl groups represented by R^9 include for example C_{1-6} alkyl groups substituted by one or more halogen atoms, or nitro, amino or hydroxy groups.

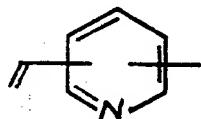
In general, in compounds of formula (1) the linker group is preferably an optionally substituted C_{1-10} alkylene, (especially C_{1-6} alkylene such as methylene, ethylene, propylene, butylene, pentylene or hexylene) C_{2-10} alkenylene or C_{2-10} alkynylene chain optionally interrupted by one or more -O- or -S- atoms or cyclohexylene, phenylene, substituted phenylene, $-NH-$, $-N(CH_3)-$, $-CONH-$, $-CONH(CH_3)-$, $-NHCO-$ or $-N(CH_3)CO-$ groups.

15 20 Particular examples of linker groups represented by L include, for example, $-(CH_2)_d-$ (where d is an integer 1 to 4 inclusive), $-(CH_2)_d-\text{C}_6\text{H}_4-$, $-(CH_2)_d-\text{C}_6\text{H}_4-\text{CH}_2\text{NHCO}-\text{C}_6\text{H}_4-\text{CH}_2-$, $-(CH_2)_d\text{NHCO}(CH_2)_e-$ (where e is an integer 1 to 4 inclusive) and $-(CH_2)_d\text{NHCO}(CH_2)_e\text{OCH}_2-$.

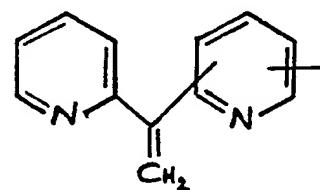
25 The reactive functional group represented by Z in compounds of formula (1) may be any group capable of reacting with a thiol, amino, carboxyl, aldehyde, aromatic or heteroaromatic group. Aromatic groups include, for example, phenolic groups. Heteroaromatic groups include for example imidazolyl groups.

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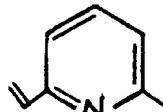
Thus, Z may be, for example, a halogen atom, for example a chlorine, bromine or iodine atom or a group selected from -SH, -NH₂, hydrazine (-NNHH₂) or a derivative thereof, [for example -N(CH₃)NH₂, -NHCONHNH₂, -NHCSNHNH₂, or phenyl hydrazine], -NCO, -NCS, -COR¹⁰, [where R¹⁰ is a halogen atom such as a chlorine or bromine atom, or a N₃, C₁₋₆ alkoxy, e.g. methoxy, C₆₋₁₂ aryloxy (e.g. nitrophenyloxy or dinitrophenyloxy), imidyloxy (e.g. succinimidylxyloxy) or imidazolyloxy group], imide, e.g. maleimide, a vinyl group of formula -Het¹-C(Het²)=CH₂ (where Het¹ and Het², which may be the same or different, is each a nitrogen containing heterocyclic group, e.g. a pyridyl group or Het¹ is a nitrogen containing heterocyclic group and Het² is a hydrogen atom) for example a vinyl pyridyl group of formula



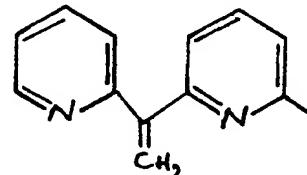
or



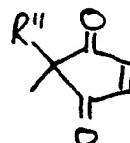
especially,



or



15 or a dione of formula

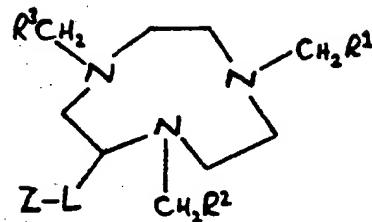
(where R¹¹ is a C₁₋₄ alkyl e.g. methyl group).

Metal complexes of the compounds of formula (1) include complexes wherein the metal is divalent or tripative and has a coordination number from 2 up to 6, especially 6. The metal may be a radioactive isotope. Examples of suitable metals include indium (In),
5 copper(Cu), lead (Pb), bismuth (Bi), cobalt (Co) and gallium (Ga). In, Ga, Co, and Cu are preferred, particularly In and Ga. ^{111}In , ^{69}Ga and ^{71}Ga are particularly preferred.

In general, optimum binding of the metal to the compounds of formula (1) may be achieved by selection of the ring size and where appropriate by adjusting the potential coordination number by choice of the group $-(\text{CH}_2)_q\text{R}^1$, $-(\text{CH}_2)_q\text{R}^2$, and/or $-(\text{CH}_2)_q\text{R}^3$. Thus a particularly important class of compound of formula (1) is that wherein p is zero. Especially useful compounds are those wherein p is zero, m is an integer 1 and n is an integer 10. 1. In general, compounds of formula (1) in which $-(\text{CH}_2)_q\text{R}^1$, $-(\text{CH}_2)_q\text{R}^2$ and $-(\text{CH}_2)_q\text{R}^3$ is each $-\text{CH}_2\text{P}(\text{O})(\text{OH})\text{R}^4$ - where R^4 is -H, $-\text{OCH}_3$, $-\text{OCH}_2\text{CH}_3$ or an alkyl group, especially a methyl group, are particularly useful.

20 Salts of the compounds of formula (1) include salts with inorganic or organic bases, for example alkali metal or alkaline earth metal salts such as lithium, sodium, potassium, magnesium or calcium salts; amine salts, such as those from primary, secondary or tertiary amines, for example ethanolamine, diethanolamine, morpholine, glucamine, N-methylglucamine or N,N-dimethylglucamine 25 salts; and amino acid salts such as lysine, arginine and ornithine salts; or acid addition salts such as hydrobromides or hydrochlorides. Pharmaceutically acceptable salts are particularly preferred.

30 An important group of compounds according to the invention has the formulae (1a):



(1a)

wherein R^1 , R^2 , R^3 , L and Z are as defined for formula (1) and metal complexes and/or salts thereof.

Compounds of this type in which R^1 , R^2 , and R^3 is each $P(O)(OH)R^4$ where R^4 is -H, $-OCH_2CH_3$ or, especially $-CH_3$ 5 are particularly preferred.

Compounds of formula (1a) in which L is a linker group [particularly those specifically identified for compounds of formula (1)] are especially useful. Particular groups of this type are those of formulae $-(CH_2)_d-$ and $-(CH_2)_dNHCO(CH_2)_e-$, where d and e, 10 which may be the same or different is each an integer 1 to 4 inclusive.

Z in compounds of formula (1a) is preferably a reactive functional group, [particularly those specifically identified for compounds of formula (1)], especially a group of formula $-NH_2$, $-COR^{10}$, 15 $-Het^1-C(Het^2)=CH_2$ or a dione of formula: $\begin{array}{c} R^u \\ | \\ \text{C}=\text{O} \\ | \\ \text{C}=\text{O} \end{array}$



Indium and gallium complexes of the compounds of formula (1a) are particularly useful.

A further group of compounds of formula (1a) which is particularly useful is that wherein L in the compounds is a covalent bond, and Z

is a hydrogen atom, and metal complexes and/or salts thereof. Gallium complexes of compounds of this type are particularly important, especially ^{71}Ga complexes, and particularly gallium complexes where R^1 , R^2 , and R^3 is each $\text{P}(\text{O})(\text{OH})\text{R}^4$,
5 especially $\text{P}(\text{O})(\text{OH})\text{CH}_3$.

The compounds of formula (1) and the metal complexes and/or salts thereof have a diagnostic use, for example as imaging agents in vitro and in vivo. The compounds of formula (1a) and the salts thereof are especially useful for use as imaging agents,
10 particularly the gallium complexes, which are particularly useful in nuclear magnetic resonance imaging, especially the gallium (particularly ^{71}Ga) complexes of formula (1a) wherein L is a covalent bond, Z is a hydrogen atom and R^1 , R^2 and R^3 is each a group $\text{P}(\text{O})(\text{OH})\text{R}^4$ especially $\text{P}(\text{O})(\text{OH})\text{CH}_3$ and the salts thereof

15 The compounds of formula (1) and the metal complexes and/or salts thereof are also cytotoxic agents and may be used in the treatment of abnormal cell disorders, for example in the treatment of tumours. For use as diagnostic and/or therapeutic agents, the compounds may be employed using conventional methods (e.g. for
20 formulation and presentation), already in use for metal complexing reagents.

For application of the compounds of formula (1) as imaging or cytotoxic agents, it may be preferable to couple the compounds to other molecules such as proteins, especially antibodies, peptides or carbohydrates to form conjugate compounds, and the compounds of formula (1) are particularly well adapted for use in this respect.
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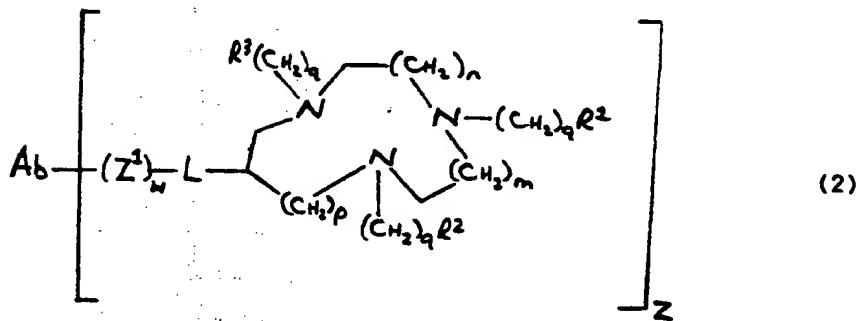
The compound of formula (1) may be coupled through any thiol, amino, carboxyl, hydroxyl, aldehyde, aromatic or heteroaromatic group present in the protein, peptide or carbohydrate.

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Thus in a further aspect of the invention, we provide a conjugate compound which comprises a compound of formula (1) or a metal complex and/or salt thereof, coupled to a protein, peptide or carbohydrate.

5 It is to be understood that a conjugate compound according to the invention may comprise more than one molecule of a compound of formula (1), coupled to any one protein, peptide or carbohydrate molecule.

10 In a particular aspect, the invention provides a conjugate compound of formula (2):



wherein m , n , p , q , R^1 , R^2 , R^3 , and L are as defined for formula (1);

15 Z^1 is the residue of a reactive functional group;
 w is zero or an integer 1;
 z is an integer 1 or more;
 Ab is an antibody; and metal complexes and/or salts thereof.

In the compound of formula (2), the residue or a reactive functional group represented by Z^1 may in general be the residue of a reactive functional group Z as defined for formula (1).

20 In particular, Z^1 may be for example $-S-$, $-NH-$, $-NHN-$, $-N(CH_3)N-$, $-NHCONHN-$, $-NHCSNHN-$, $-N(Ph)N-$ (where Ph is phenyl), $-NC(O)-$, $-NC(S)-$, $-CO-$, $-N$, $-H^1-C(Het^2)CH_2-$ or R^u

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The antibody in the conjugate formula (2) may in general belong to any immunoglobulin class. Thus for example it may be an immunoglobulin M antibody or, in particular, an immunoglobulin G antibody. The antibody molecule may be of animal, for example 5 mammalian origin, and may be for example of murine, rat or human origin. It may be a natural antibody or a fragment thereof, or, if desired, a recombinant antibody or antibody fragment i.e. an antibody molecule or antibody fragment which has been produced using recombinant DNA techniques.

10 Particular recombinant antibodies or antibody fragments include, (1) those having an antigen binding site at least part of which is derived from a different antibody, for example those in which the hypervariable or complementarity determining regions of one antibody have been grafted into the variable framework regions of a second, 15 different antibody (as described in European Patent Specification No. 239400); (2) recombinant antibodies or fragments wherein non-Fv sequences have been substituted by non-Fv sequences from other, different antibodies (as described in European Patent Specification Nos. 171496, 173494 and 194276; or (3) recombinant antibodies or 20 fragments possessing substantially the structure of a natural immunoglobulin but wherein the hinge region has a different number of cysteine residues from that found in the natural immunoglobulin, or wherein one or more cysteine residues in a surface pocket of the recombinant antibody or fragment is in the place of another amino acid residue present in the natural immunoglobulin (as described in 25 International Patent Applications Nos. PCT/GB 88/00730 and PCT/GB 88/00729 respectively).

The antibody may be of polyclonal or, preferably, monoclonal origin. It may be specific for any number of antigenic 30 determinants, but is preferably specific for one. The antigenic determinants may be any hapten or antigenic determinants associated with any antigen. Particular antigens include those associated with

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5 animals, e.g. humans, [for example normal animal tissue or organ
10 cell-associated antigens, tumour cell-associated antigens (for
example oncofetal antigens such as carcinoembryonic antigen or
15 alphafetoprotein, placental antigens such as chorionic gonadotropin
and placental alkaline phosphatase, and prostate antigens such as
prostatic acid phosphatase and prostate specific antigen) and
antigens associated with components of body fluids such as fibrin or
20 platelets], viruses, bacteria and fungi.

25 In a preferred aspect the antibody may be capable of recognising and
binding a tumour cell-associated antigen, particularly one or more
epitopes on the TAG-72 antigen associated with human breast and
colon tumours. A particularly preferred antibody of this type is
the monoclonal antibody B72.3 [Colcher, D. et al Proc. Nat. Acad.
Sci. USA (1981), 78 3199] or a fragment thereof, particularly a
15 $F(ab')_2$ fragment.

20 The antibody Ab will in general be coupled to the remainder of the
conjugate of formula (2) (i.e. the macrocycle and linker) through
any appropriate reactive atom or group, for example a nitrogen or,
especially, sulphur atom, present in the antibody. It will be
appreciated that any one antibody molecule may contain more than one
reactive group capable of coupling with the macrocycle and linker.
Thus, for example, z in the conjugates of formula (2) may be an
integer 1, 2, 3, 4, 5, 6 or more depending on the number of
25 macrocycles linked to any particular antibody molecule or fragment.

30 Indium and gallium complexes of conjugates of formula (2) are
particularly useful.

It is to be understood that the definitions and preferences
expressed for m, n, p, q, R^1 , R^2 , R^3 and L in compounds of
formula (1), and for classes of compounds of formula (1) are also
30 applicable to conjugates of formula (2).

Particularly useful conjugate compounds according to the invention are those comprising a compound of formula (1a), or a metal complex and/or salt thereof, coupled to an antibody. The indium and gallium complexes of these conjugates are especially important.

5 The compounds of formulae (1) and (2) may be formulated for use in accordance with conventional practice, and thus according to a further respect of the invention we provide a composition comprising a compound of formula (1) or a compound of formula (2) or a metal complex and/or salt thereof, together with one or more 10 pharmaceutically acceptable carriers.

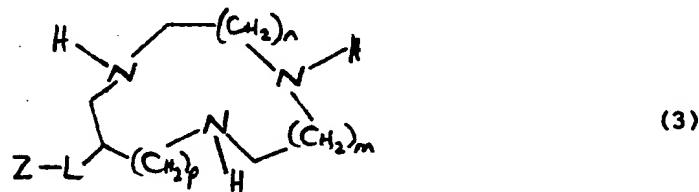
Particularly suitable compositions according to the invention are those adapted for parenteral administration, especially intravenous administration. Suitable formulations of this type include solutions of the compounds of formulae (1) or (2) in isotonic saline.

15 The quantities of compounds of formulae (1) or (2) used in formulations according to the invention will vary according to the intended use (i.e. imaging or therapy) and other variables such as the intended cell target, but may be easily determined in accordance with conventional practice for reagents of this type.

20 Compounds of the invention may be prepared by the following processes wherein the groups and symbols R^1 , R^2 , R^3 , m, n, p, q, L, Z, Ab and z are as defined for formulae (1) and (2) except where stated otherwise. Where a metal complex is desired as a final product, the complexation with a metal atom may be carried out as a 25 final step in the production process, as described below for the complexation of compounds of formula (1), or alternatively it may be desirable to complex the metal at an earlier stage in the process, providing of course that the requisite macrocycle structure is present. In the following processes, it may be desirable to use starting materials in which the group Z is in a protected state, or 30 which contain a precursor of the group, as discussed below.

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Thus, according to a further aspect of the invention a compound of formula (1) [wherein q is an integer 1-6 and, where present, in R¹, R² and/or R³ the group X is an oxygen atom] or a metal complex thereof may be prepared by reaction of a corresponding compound of formula (3)



or a metal complex thereof, with a reagent

10 D(CH₂)_qP(O)(OR¹³)⁴, or D(CH₂)_qP(O)(OR¹³)₂, (where D is a displaceable group, for example a halogen atom such as a bromine atom; R¹³ is a C₁₋₄ alkyl, e.g. methyl or ethyl group; and q and R⁴ are as defined previously) followed where necessary by hydrolysis.

15 The reaction may be performed in a solvent such as water or an organic solvent such as a nitrile e.g. acetonitrile or an alcohol, e.g. ethanol or an amide e.g. dimethylformamide in the presence of a base such as an alkali metal carbonate or hydroxide, e.g. sodium, potassium or caesium carbonate, or sodium, potassium or lithium hydroxide, at an elevated temperature e.g. the reflux temperature.

20 Where appropriate, hydrolysis may be achieved using a base, such as described above, in a suitable solvent, for example sodium hydroxide in an alcohol such as ethanol.

In this reaction, the group Z may need to be in a protected state. Conventional protecting groups may be used, depending on the nature

of Z, and may be removed using standard procedures, once the desired reaction has been effected.

Reagents $D(CH_2)_q P(O)(OR^{13})_2^4$ and $D(CH_2)_q P(O)(OR^{13})_2^2$ may be prepared by heating compounds of formulae $P(OR^{13})_2^2 R^4$ or 5 $P(OR^{13})_3^3$ with a compound $(CH_2)_q D_2$.

In an alternative process, a compound of formula (1) [wherein in R¹, R² and/or R³ the group X where present is an oxygen atom] may be prepared by reaction of a compound of formula (3), or a metal 10 complex thereof with a phosphine $R^4 P(OR^{13})$ in the presence of suitable aldehyde (for example formaldehyde or paraformaldehyde), followed by hydrolysis.

The reaction may be performed in an organic solvent, e.g. a nitrile, alcohol, or amide, or an ether such as tetrahydrofuran at an elevated temperature, for example the reflux temperature.

15 Hydrolysis may be achieved using an acid, for example an inorganic acid such as hydrochloric acid, at an elevated temperature such as the reflux temperature.

Compounds of formula (1) may also be prepared by interconversion 20 from other compounds of formula (1). Thus one functional group Z may be exchanged for another and, if desired a linker group L changed to another by appropriate manipulative reactions. For example, a compound of formula (1) where -L-Z is a group $-L^1-NHCO-L^2-Z$ (where $-L^1-NHCO-L^2$ represents the group L) may be prepared by reaction of a corresponding compound wherein -L-Z 25 represents $-L^1-NH_2^b$ with a reagent R^b-O-L^2-Z (where R^b is for example an amide, such as succinimide, or a substituted phenyl group such as a p-nitrophenyl group) in the presence of a tertiary amine, such as diisopropylethylamine, in a solvent such as dimethylformamide.

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Reagents of formula R^bO-L^2-Z are either known compounds or may be obtained from known starting materials using methods analogous to those used for the preparation of the known compounds.

5 In another interconversion process, a compound of formula (1) wherein X where present is a sulphur atom may be prepared by reaction of a corresponding compound wherein X is an oxygen atom by reaction with a sulphide, for example phosphorous pentasulphide, at an elevated temperature.

10 It will be appreciated that where it is desired to prepare a compound of formula (1) in which one or two of R^1 , R^2 and R^3 are a hydrogen atom or a CO_2H group this may be achieved by first selectively N-protecting the compound of formula (3) or a precursor using an appropriate amine protecting group(s), for example a p-toluenesulphonyl group in accordance with conventional practice.

15 Reaction of the N-protected compound (3) to introduce the required group $-P(O)(XH)R^4$ using the methods described above followed by deprotection and further reaction with a group $D(CH_2)_qH$ or $D(CH_2)_qCO_2H$ (or an acid protected form thereof) using the reagents and conditions described previously for the introduction of 20 the group $-P(O)(XH)R^4$ then yields the desired compound in which one or two of R^1 , R^2 and R^3 are -H or CO_2H .

25 Where metal complexes of compounds of formulae (1) or (2) are required (or any other suitable macrocyclic intermediate described herein) these may be prepared by treating the compound with a metal salt (for example a metal halide e.g. a chloride, or a nitrate, acetate, carbonate or sulphate) or a metal oxide in an appropriate solvent for example an aqueous or non aqueous solvent, (e.g. acetonitrile, acetone, propylene carbonate, dimethylformamide or dimethylsulphoxide) at any suitable temperature from $0^{\circ}C$ to 30 $100^{\circ}C$ such as $10^{\circ}C$ to $80^{\circ}C$ e.g. around $60^{\circ}C$.

A conjugate compound of formula (2) or a metal complex thereof may be prepared by reaction of a corresponding compound of formula (1) or a metal complex thereof with an antibody Ab (as previously defined).

5 The reaction may be performed in a suitable solvent, for example an aqueous solvent such as a phosphate buffer, at an appropriate temperature, for example at 0°C-30°C, especially 0°-10°C e.g. 4°C.

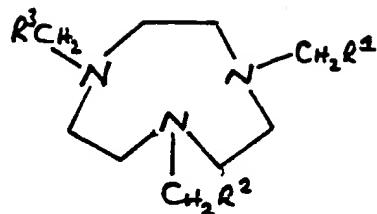
10 The antibody Ab may be obtained using procedures well known in the art. If desired, before the coupling reaction, the antibody may first be treated to yield appropriate groups for reaction with the compound of formula (1). Thus for example the antibody may be subjected to oxidation, for example periodate oxidation to yield aldehyde groups, or, in particular, may be treated with a reagent [e.g. Traut's reagent (2-iminothiolane)] using standard procedures 15 to generate free sulphhydryl groups in the molecule.

20 Salts of compounds of formulae (1) or (2) and their metal complexes may be prepared by conventional means, for example by reaction of a compound of formulae (1) or (2) with an appropriate base or acid in a suitable aqueous or non-aqueous solvent as described above, at any suitable temperature from 0°C to 100°C.

Intermediates of formula (3) may be prepared by the methods described in International Patent Specification Publication No. WO 89/01475.

The invention is illustrated by the following Examples:

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Example 1

(a) R^1, R^2, R^3 is each $P(O)(OCH_2CH_3)CH_3$
 (b) R^1, R^2, R^3 is each $P(O)(OH)CH_3$
 (c) Indium complex of (b)
 5 (d) Gallium complex of (b)

(a) To a solution of 1,4,7-triazacyclononane (90mg) and paraformaldehyde (0.10g) in dry tetrahydrofuran (5ml) was added methyl diethoxyphosphine (0.5g) and the solution was boiled under reflux (under N_2) for 48h. After removal of solvent, the residue was chromatographed on neutral alumina (eluent 0 \rightarrow 10 5% methanol/CH₂Cl₂) to yield the desired triester product as a pale-yellow oil, R_f = 0.51 (3% methanol- CH₂Cl₂). δp (CD₂Cl₂) 54.2. m/e (DCI, CH₂Cl₂) 491, 490 [M⁺+1] 382, 289.

15 (b) The triester from (a)(50mg) was treated with hydrochloric acid (6M, 2ml) and was heated to 110° for 18h. Removal of solvent yielded the desired triacid as a glassy solid m/e (FAB, m-nitrobenzylalcohol) 406 (M⁺+1). δp (H₂O, pH1) 42.5.

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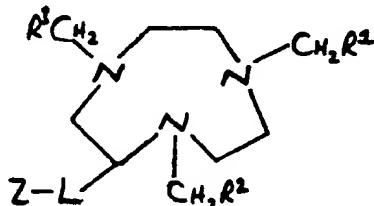
δ_H (D_2O) 3.32 (18H, mult., CH_2CH_2N , + CH_2P), 1.42 (9H, d, $J=14$, $P-CH_3$).

5 Admixture of equimolar quantities of the triacid prepared in (b) and either a solution of indium nitrate in dilute nitric acid (pH1) or gallium nitrate in nitric acid (pH1) yielded solutions of the corresponding complexes quantitatively with the following characteristics:

(c) Indium complex : δp (H_2O , pH1) + 38.3 - invariant over 4 weeks at pH1

10 (d) Gallium complex : δp (H_2O , pH1) + 40.5; $\delta Ga(H_2O$, pH1) +137 ppm (invariant pH0 for 1 week); m/e (DCI methanol) = 474, 472 ($M^+ + 1$).

Example 2



15 (a) R^1 , R^2 , R^3 is each $P(O)(OCH_2CH_3)CH_3$; L-Z is $-(CH_2)_4NHCO$ -phenyl

(b) R^1 , R^2 , R^3 is each $P(O)(OH)CH_3$; L-Z is $-(CH_2)_4NH_2$.

(c) R^1 , R^2 , R^3 is each $P(O)(OH)CH_3$; L-Z is $-(CH_2)_4NHCO(CH_2)_2CO_2$ -Ph where Ph is 4-nitrophenyl.

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(a) To a solution of 2-(4-N-benzamidyl)butyl-1,4,7,-triazacyclononane [prepared as described in International Patent Specification No. WO89/01475] (0.39g) in dry tetrahydrofuran (10ml) was added methyldiethoxyphosphine (0.66g) and paraformaldehyde (0.18g) and the mixture was heated to reflux for 18h with removal of water (Soxhlet: 4A molecular sieves). After filtration and removal of solvent, the residue was purified by chromatography on neutral alumina (0 → 3% methanol in CH_2Cl_2) to yield the desired tri-ester product as a colourless oil (0.26g). m/e (d.c.i.) 665 ($\text{M}^+ + 1$), 557, 437, 126, 109. δ_{C} (CDCl_3) 167.4 (carbonyl); 134.75, 130.67, 121.86, 127.10 (aryl); 63.48 (CH_2O); 59.87, 59.03, 58.2, 58.15, 57.9, 57.8, 57.0, 53.1, 53.0, 52.7, (CH_2N); 39.4, 39.35 (CH_2NHCO , diastereoisomers); 28.6 (br.s, CH_2C); 24.0, 23.9 (CH_2C); 16.40, 16.36 ($\text{CH}_3\text{CH}_2\text{O}$: diastereomers); 13.22 (d, $J_{\text{CP}} = 89\text{Hz}$, major), 12.95, 12.80 (d+d, $J_{\text{CP}} = 90$, 91Hz, CH_3P minor diastereoisomers) ν_{max} (film) 3405, 3200 (NH); 2960, 2915, 2825 (CH); 1640 (NHCO, s); 1205 (vs); 1035 (vs), 950 (s) cm^{-1} . δ_{H} (CDCl_3) 7.91 (2H, dd, ortho arom.), 7.85 (1H, br. t, NHCO), 7.45-7.39 (3H, para + meta arom CH), 4.02 (6H, mult, CH_2O), 3.50-2.60 (19H, mult, $\text{CHN} + \text{CH}_2\text{N}$) 1.63 (2H, mult, $\text{CH}_2\text{CH}_2\text{NHCO}$) 1.55-1.42 (4H, mult, CH_2C), 1.30 (9H, mult, CH_3P), 1.20 (9H, t+t+t, $\text{CH}_3\text{CH}_2\text{O}$, diastereomers)

(b) A solution of the triester (0.13g), prepared in (a) in hydrochloric acid (6M, 5ml) was heated to reflux for 24h. After cooling, washing with ether (3 x 2ml), dichloromethane (2 x 5ml) and evaporation under high vacuum (0.01mm Hg, 18h), a colourless glass was obtained of the tetrahydrochloride salt of the corresponding triacid in which L-Z is $(\text{CH}_2)_4\text{NH}_2$. m/e (F.a.b.; glycerol): 477 (M^+).

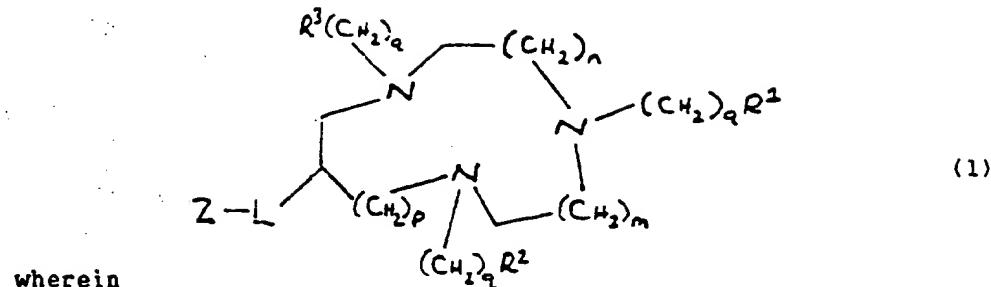
- 21 -

(c) T a s luti n of the triacid prepared in (a) (0.048g) in dry DMSO (0.5ml) was add d N-methylmorph line (90mg) and di-(p-nitrophenyl)-succinate (54mg) and the mixture was stirred for 18h at 20°^oC, monitoring by HPLC (Dynamax C18 60A, 21.4mm column: A=0.1% trifluoroacetic acid -H₂O), C=0.1% trifluoroacetic acid -CH₃CN: t=0 A=95%, C=5%; t=20 min A=5%, C=95%; flow= 10ml min⁻¹) R_t=12 mm. Purification by HPLC afforded the desired tri-acid as a colourless solid (25mg). m/e (F.a.b., glycerol) 699 (M⁺). δ_H(D₂O) 8.30 (2H, 5
ortho Ar), 7.34 (2H, ortho Ar), 4.0 2.5 (23H, mult, 10
CH₂CO+CH₂N), 1.8-1.5 (6H, mult, CH₂C), 1.25 (d+d+d, 9H, 15
CH₃P).

Reaction of the compound of part (c) with gallium nitrate or indium nitrate as described in Example 1 yielded the 15 corresponding gallium or indium complexes.

CLAIMS

1. A compound of formula (1):



m and n, which may be the same or different, is each zero or an integer 1, 2 or 3;

5 p is zero or an integer 1 or 2;

q is zero or an integer from 1 to 6 inclusive;

10 R¹, R² and R³, which may be the same or different, is each a hydrogen atom or a group -CO₂H or -P(O)(XH)R⁴ (where X is an oxygen or sulphur atom and R⁴ is a hydrogen atom or an alkyl or alkoxy group), with the proviso that at least one of R¹, R² and R³ is a -P(O)(XH)R⁴ group;

L is a covalent bond or a linker group;

Z is a hydrogen atom or a reactive functional group;

15 and metal complexes and/or salts thereof.

2. A compound according to Claim 1 where R¹, R² and R³ are the same and is each a group -P(O)(XH)R⁴.

3. A compound according to Claim 2 wherein R¹, R² and R³ is each a group -P(O)(OH)CH₃.

20 4. A compound according to any of the preceding claims wherein q is an integer 1.

5. A compound according to any of the preceding claims wherein L is a covalent bond and Z is a hydrogen atom.

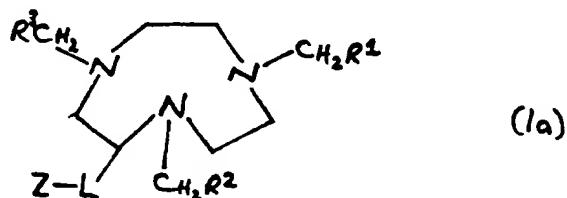
- 23 -

6. A compound according to any of Claims 1 to 4 wherein L is an optionally substituted aliphatic hydrocarbyl chain, optionally interrupted by one or more heteroatoms selected from -O- or -S- or by one or more $-N(R^5)_2$, $-N(R^5)CO-$, cycloaliphatic, aromatic, or heteroaromatic groups.

5

7. A compound according to any of Claims 1 to 4 and 6 wherein Z is any group capable of reacting with a thiol, amino, carboxyl, aldehyde, aromatic or heteroaromatic group.

8. A compound of formula (1a):



10 wherein
 R^1 , R^2 , and R^3 , which may be the same or different, is each a hydrogen atom or a group $-CO_2H$ or $-P(O)(XH)R^4$ (where X is an oxygen or sulphur atom and R^4 is a hydrogen atom or an alkyl or alkoxy group), with the proviso that at least one of R^1 , R^2 and R^3 is a $-P(O)(XH)R^4$ group;

15 L is a covalent bond or a linker group;

Z is a hydrogen atom or a reactive functional group; and metal complexes and/or salts thereof.

9. A compound according to Claim 8 wherein R^1 , R^2 and R^3 is each a group $-P(O)(OH)CH_3$.

20

10. A compound according to Claims 8 or 9 wherein q is an integer 1.

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11. A compound according to Claims 8 to 10 wherein L is an
5 partially substituted aliphatic hydrocarbyl chain, partially
interrupted by one or more heteroatoms selected from -O- or -S-
or by one or more -N(R⁵)-, -N(R⁵)CO-, cycloaliphatic,
aromatic, or heteroaromatic groups and Z is any group capable
of reacting with a thiol, amino carboxyl, aldehyde, aromatic or
heteroaromatic group.
12. A compound according to Claims 8-10 wherein L is a covalent
bond and Z is a hydrogen atom.
- 10 13. An indium or gallium complex of any of the preceding claims.
14. A gallium complex of a compound according to Claim 12.

INTERNATIONAL SEARCH REPORT

International Application No PCT/GB 90/00982

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC: 5 C 07 F 9/6515, A 61 K 49/02, A 61 K 43/00, A 61 K 39/395

II. FIELDS SEARCHED

Minimum Documentation Searched ⁷

Classification System ⁸	Classification Symbols
IPC ⁵	C 07 F 9/00

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched ⁹

III. DOCUMENTS CONSIDERED TO BE RELEVANT¹⁰

Category ¹¹	Classification of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	Bulletin of the Academy of Sciences of the USSR, Division of Chemical Science, vol. 33, no. 4, part 1, April 1984, Plenum Publishing Corporation, (New York, US), M.I. Kabachnik et al.: "Synthesis and study of a new complexone - N, N',N"-Tris-(Dihydroxyphosphorylmethyl)-1,4,7-triazacyclononane", pages 769-777, see the whole article --	1,4,5,8,10, 12
A	WO, A, 89/01475 (CELLTECH LIMITED) 23 February 1989 see the whole document (cited in the application) --	1-14
		. / .

* Special categories of cited documents: ¹⁰

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"Z" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search
20th November 1990

Date of Mailing of this International Search Report

29.12.90

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

N. TORIBIA *Nicola TORIBIA*

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages	Relevant to Claim No.
A	WO, A, 86/02352 (BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM) 24 April 1986 see the whole document (cited in the application) --	1, 4, 5, 8, 10, 12
A	EP, A, 0197437 (KONISHIROKU PHOTO INDUSTRY CO., LTD) 15 October 1986 see pages 10-15; claims (cited in the application)	1, 4, 5, 8, 10, 12

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

GB 9000982

SA 38227

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 11/12/90. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

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		EP-A-	0329737	30-08-89
		JP-T-	2501069	12-04-90
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